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(FILE 'HOME' ENTERED AT 09:35:28 ON 24 AUG 2006)

FILE 'REGISTRY' ENTERED AT 09:35:48 ON 24 AUG 2006

L1 STRUC
L2 50 S L1
L3 18324 S L1 FUL
L4 2850 S L3 AND (PIPERIDIN? OR AZEPIDIN?)
E AZEPIDIN
L5 74 S L3 AND AZEPIN?

FILE 'CAPLUS' ENTERED AT 09:38:59 ON 24 AUG 2006

L6 16 S L5

FILE 'REGISTRY' ENTERED AT 09:40:41 ON 24 AUG 2006

FILE 'CAPLUS' ENTERED AT 09:42:12 ON 24 AUG 2006

FILE 'STNGUIDE' ENTERED AT 09:48:08 ON 24 AUG 2006

FILE 'CAPLUS' ENTERED AT 09:49:53 ON 24 AUG 2006

FILE 'REGISTRY' ENTERED AT 09:50:50 ON 24 AUG 2006

L7 2850 S L4 NOT L5
L8 STRUC
L9 546 SEARCH L8 SSS SUB=L7 FUL
L10 546 SEARCH L8 SSS SUB=L9 FUL

FILE 'CAPLUS' ENTERED AT 09:57:41 ON 24 AUG 2006

L11 2 S L10

FILE 'STNGUIDE' ENTERED AT 09:59:22 ON 24 AUG 2006

FILE 'REGISTRY' ENTERED AT 10:06:36 ON 24 AUG 2006

L12 2304 S L7 NOT L10

FILE 'CAPLUS' ENTERED AT 10:08:33 ON 24 AUG 2006

L13 159 S L12
L14 3 S L13 AND LIPASE
L15 1 S L13 AND PHOSPHOLIPASE

=> s l14 not l15

L16 3 L14 NOT L15

=> s l13 and (phospho?(l)lipase)

1072064 PHOSPHO?

47325 LIPASE

3940 PHOSPHO?(L)LIPASE

L17 0 L13 AND (PHOSPHO?(L)LIPASE)

=> d bib hitstr l14 1-3

L14 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:60473 CAPLUS

DN 140:128423

TI Preparation of heterocyclylbenzoylureas for treating type 2 diabetes

IN Schoenafinger, Karl; Defossa, Elisabeth; Kadereit, Dieter; Von Roedern, Erich; Klabunde, Thomas; Burger, Hans-Joerg; Herling, Andreas; Wendt, Karl-Ulrich

PA Aventis Pharma Deutschland GmbH, Germany

SO PCT Int. Appl., 67 pp.

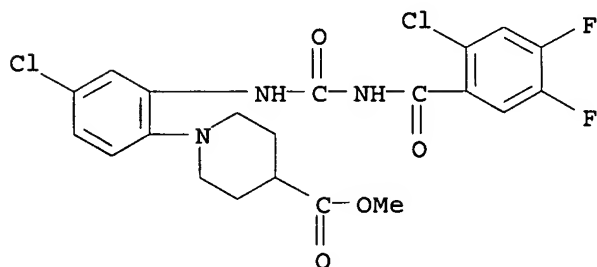
CODEN: PIXXD2

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004007455	A1	20040122	WO 2003-EP7078	20030703
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	DE 10231627	B4	20060309		
	DE 10306503	A1	20040826	DE 2003-10306503	20030217
	DE 10320326	A1	20041202	DE 2003-10320326	20030506
	CA 2493374	AA	20040122	CA 2003-2493374	20030703
	AU 2003249937	A1	20040202	AU 2003-249937	20030703
	EP 1523475	A1	20050420	EP 2003-763692	20030703
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
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	CN 1668593	A	20050914	CN 2003-816581	20030703
	JP 2006502247	T2	20060119	JP 2005-505058	20030703
	US 2004152743	A1	20040805	US 2003-617498	20030711
	ZA 2004010133	A	20051123	ZA 2004-10133	20041215
	NO 2005000648	A	20050207	NO 2005-648	20050207
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	DE 2003-10306503	A	20030217		
	DE 2003-10320326	A	20030506		
	US 2002-430782P	P	20021204		
	WO 2003-EP7078	W	20030703		
OS	MARPAT 140:128423				
IT	648917-24-2P				
	RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)				
	(preparation of heterocyclylbenzoylureas for treating type 2 diabetes)				
RN	648917-24-2 CAPLUS				
CN	4-Piperidinecarboxylic acid, 1-[4-chloro-2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]phenyl]-, methyl ester (9CI) (CA INDEX NAME)				



IT 648917-12-8P 648917-13-9P 648917-14-0P
 648917-15-1P 648917-16-2P 648917-17-3P
 648917-18-4P 648917-19-5P 648917-20-8P
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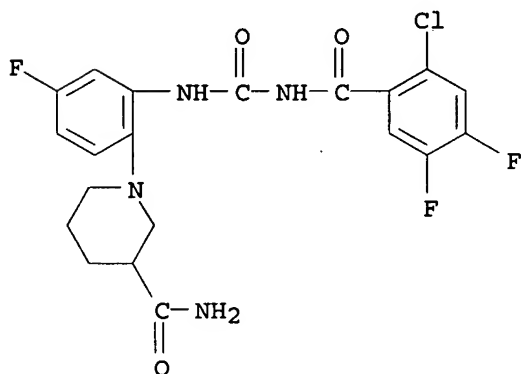
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 648917-44-6P 648917-45-7P 648917-46-8P
 648917-47-9P 648917-48-0P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
 (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
 (Uses)

(preparation of heterocyclylbenzoylureas for treating type 2 diabetes)

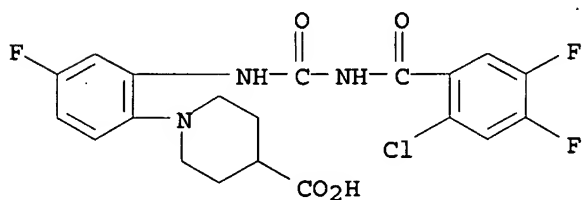
RN 648917-12-8 CAPLUS

CN 3-Piperidinecarboxamide, 1-[2-[[[(2-chloro-4,5-
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 NAME)



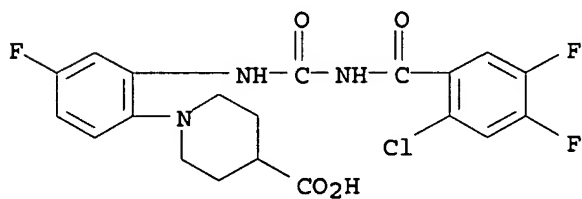
RN 648917-13-9 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-
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 NAME)



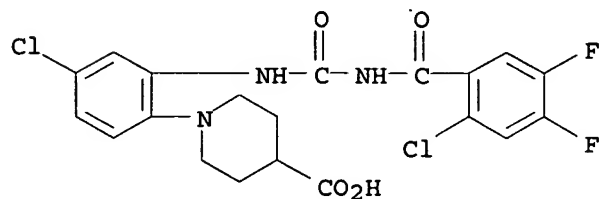
RN 648917-14-0 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-
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 (9CI) (CA INDEX NAME)



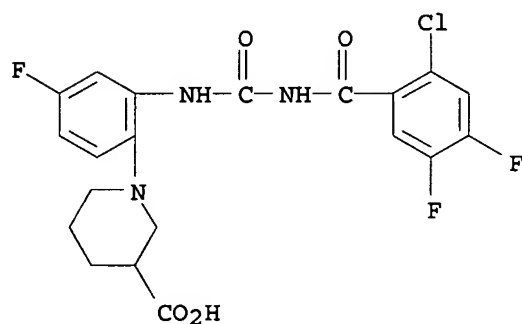
RN 648917-15-1 CAPLUS

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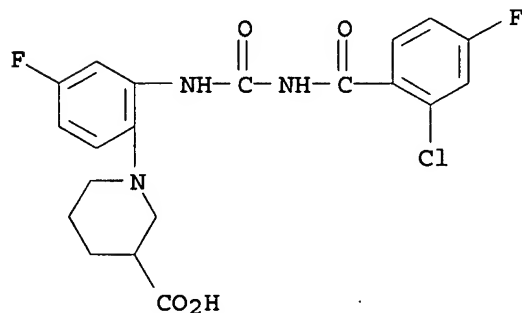
RN 648917-16-2 CAPLUS

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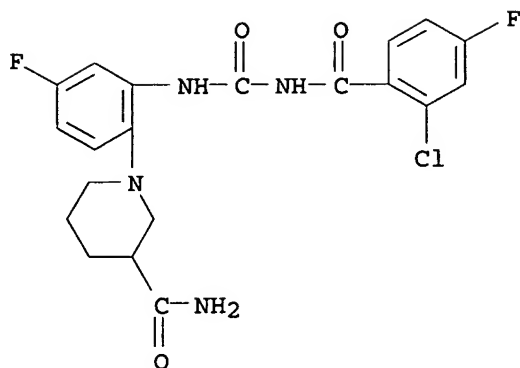
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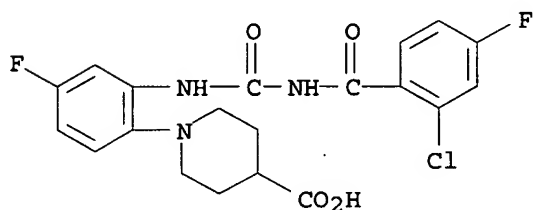
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CN 3-Piperidinecarboxamide, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]- (9CI) (CA INDEX NAME)



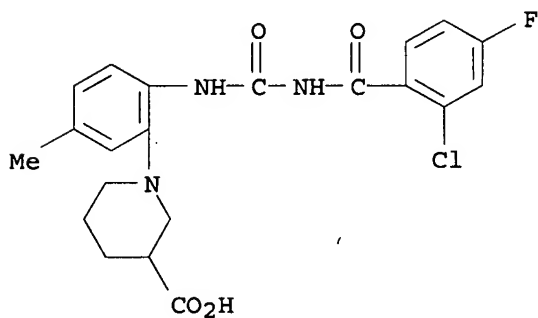
RN 648917-19-5 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]- (9CI) (CA INDEX NAME)



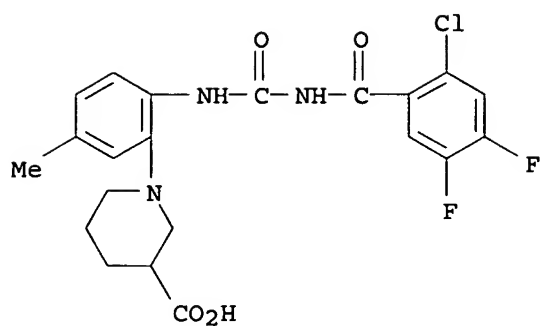
RN 648917-20-8 CAPLUS

CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-5-methylphenyl]- (9CI) (CA INDEX NAME)



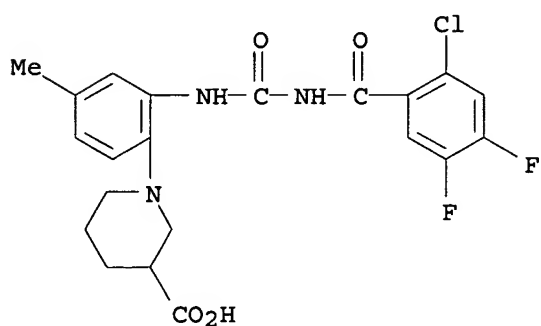
RN 648917-21-9 CAPLUS

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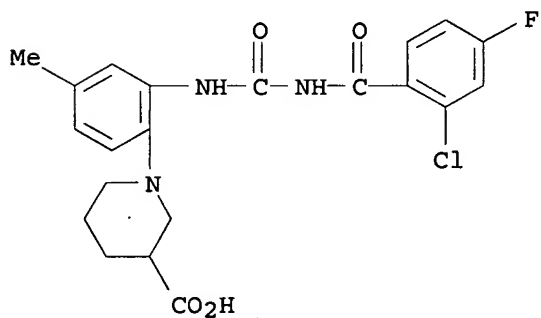
RN 648917-22-0 CAPLUS

CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-methylphenyl]- (9CI) (CA INDEX NAME)



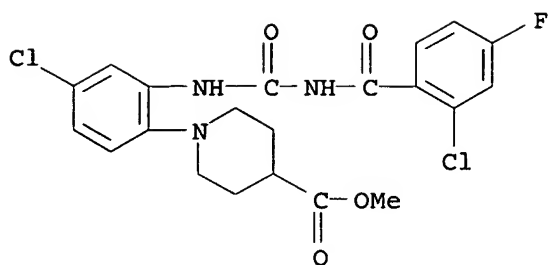
RN 648917-23-1 CAPLUS

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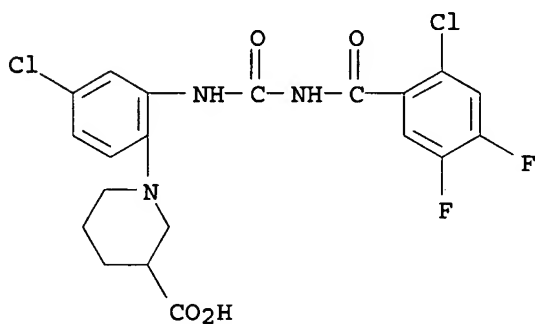
RN 648917-25-3 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[4-chloro-2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]phenyl]-, methyl ester (9CI) (CA INDEX NAME)



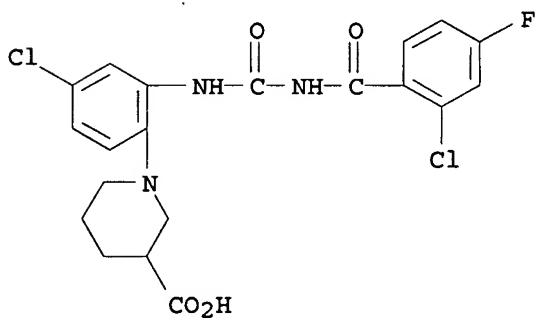
RN 648917-26-4 CAPLUS

CN 3-Piperidinecarboxylic acid, 1-[4-chloro-2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



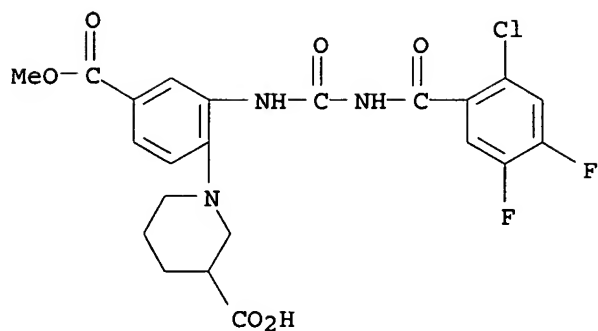
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CN 3-Piperidinecarboxylic acid, 1-[4-chloro-2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



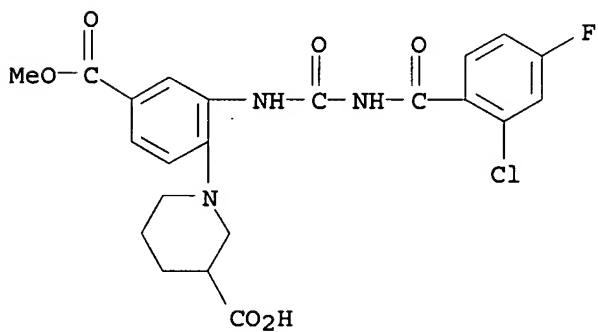
RN 648917-28-6 CAPLUS

CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-(methoxycarbonyl)phenyl]- (9CI) (CA INDEX NAME)



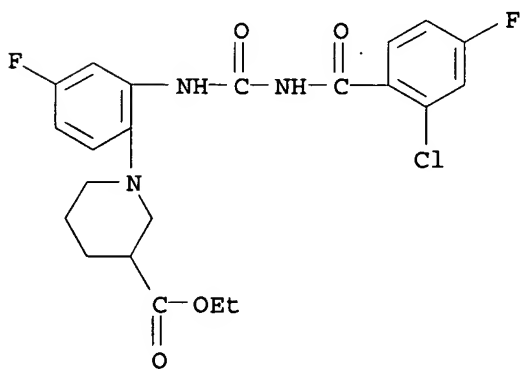
RN 648917-29-7 CAPLUS

CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-(methoxycarbonyl)phenyl]- (9CI) (CA INDEX NAME)



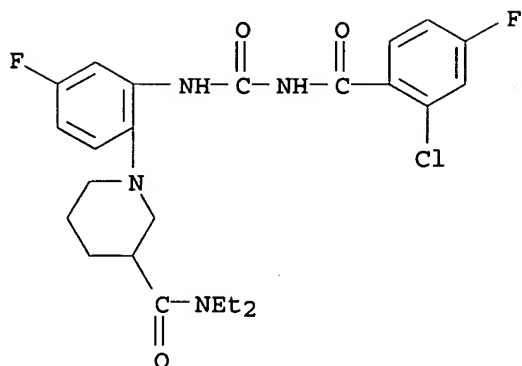
RN 648917-30-0 CAPLUS

CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-, ethyl ester (9CI) (CA INDEX NAME)



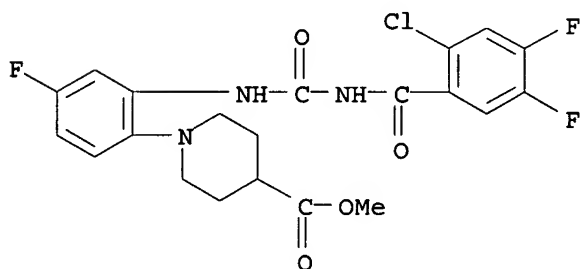
RN 648917-31-1 CAPLUS

CN 3-Piperidinecarboxamide, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-N,N-diethyl- (9CI) (CA INDEX NAME)



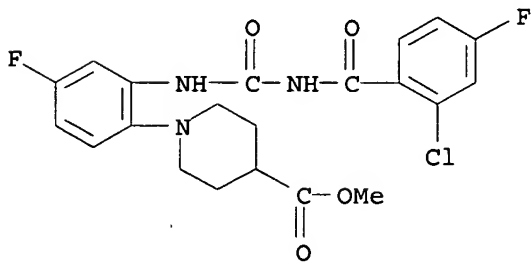
RN 648917-32-2 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-, methyl ester (9CI)
(CA INDEX NAME)



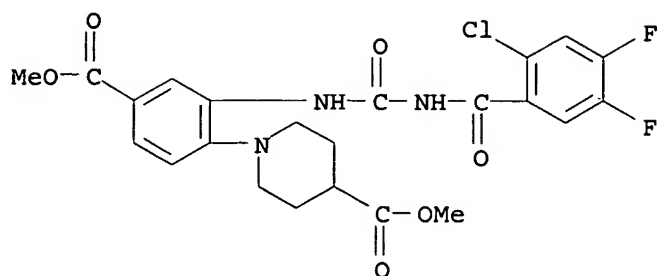
RN 648917-33-3 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-, methyl ester (9CI)
(CA INDEX NAME)



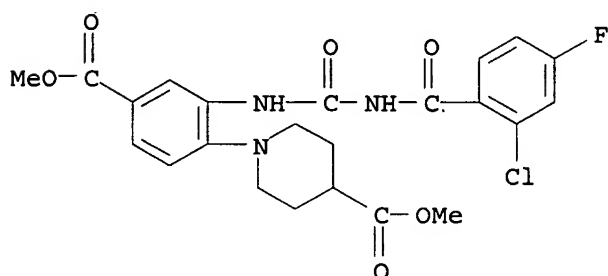
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CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-(methoxycarbonyl)phenyl]-, methyl ester (9CI) (CA INDEX NAME)



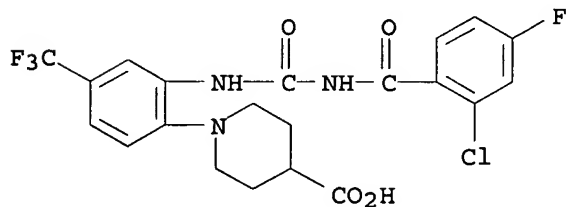
RN 648917-35-5 CAPLUS

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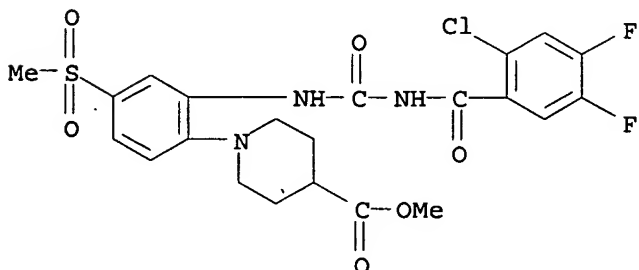
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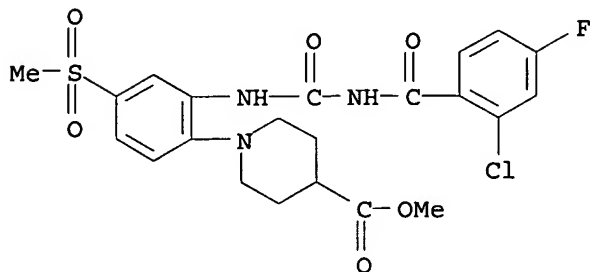
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CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-(methylsulfonyl)phenyl]-, methyl ester (9CI) (CA INDEX NAME)



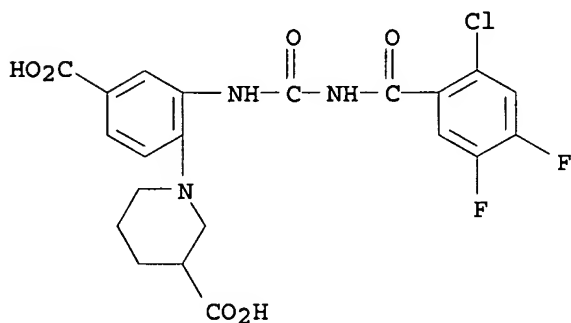
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CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-(methylsulfonyl)phenyl]-, methyl ester (9CI) (CA INDEX NAME)



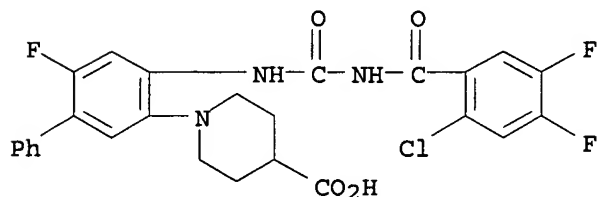
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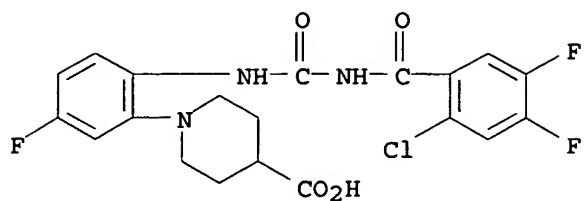
RN 648917-40-2 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[4-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-6-fluoro[1,1'-biphenyl]-3-yl]- (9CI) (CA INDEX NAME)



RN 648917-42-4 CAPLUS

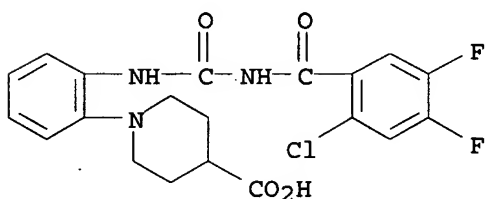
CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-5-fluorophenyl]-, monosodium salt (9CI) (CA INDEX NAME)



● Na

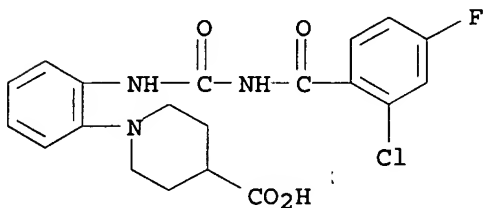
RN 648917-43-5 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 648917-44-6 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



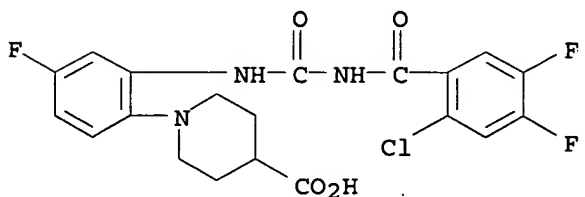
RN 648917-45-7 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-, compd. with 2-amino-2-(hydroxymethyl)-1,3-propanediol (1:1) (9CI) (CA INDEX NAME)

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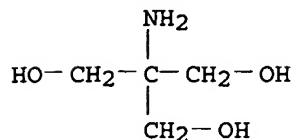
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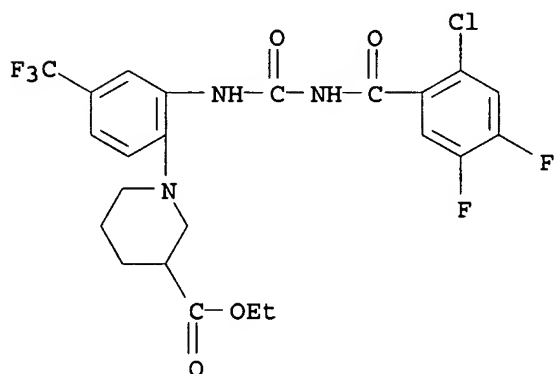


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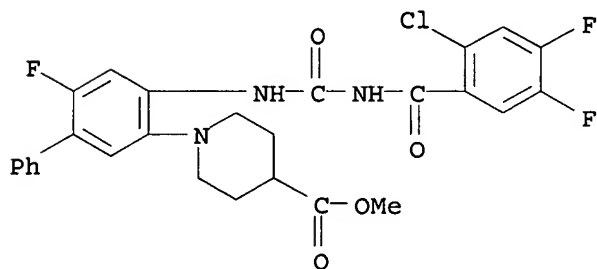
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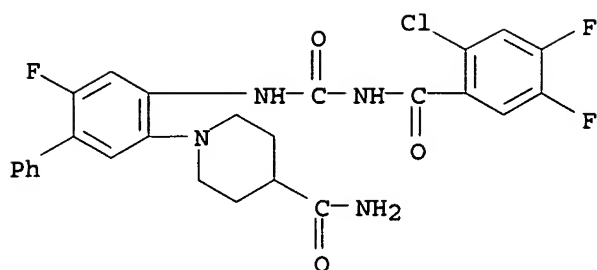
RN 648917-46-8 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl) amino] carbonyl] amino]-4-(trifluoromethyl)phenyl]-, ethyl ester (9CI) (CA INDEX NAME)



RN 648917-47-9 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[4-[[[(2-chloro-4,5-difluorobenzoyl) amino] carbonyl] amino]-6-fluoro[1,1'-biphenyl]-3-yl]-, methyl ester (9CI) (CA INDEX NAME)



RN 648917-48-0 CAPLUS
CN 4-Piperidinecarboxamide, 1-[4-[[[(2-chloro-4,5-difluorobenzoyl) amino] carbonyl] amino]-6-fluoro[1,1'-biphenyl]-3-yl]- (9CI) (CA INDEX NAME)

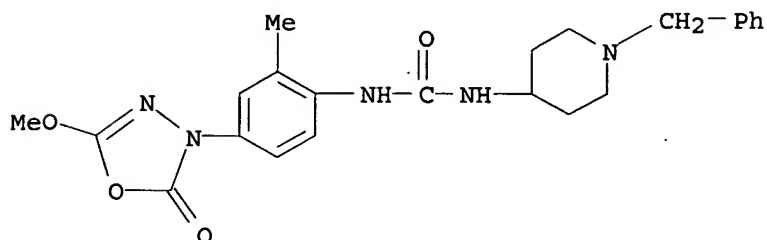


RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2003:696732 CAPLUS
DN 139:214471
TI Preparation of 5-alkoxy-3-phenyl-1,3,4-oxadiazol-2(3H)-ones for producing medicaments inhibiting pancreatic lipase
IN Schoenafinger, Karl; Petry, Stefan; Mueller, Guenter; Bauer, Armin; Heuer, Hubert Otto
PA Aventis Pharma Deutschland G.m.b.H., Germany
SO PCT Int. Appl., 63 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003072098	A1	20030904	WO 2003-EP1560	20030217
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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CA 2477005	AA	20030904	CA 2003-2477005	20030217
AU 2003210292	A1	20030909	AU 2003-210292	20030217
EP 1482929	A1	20041208	EP 2003-742942	20030217
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003008045	A	20041221	BR 2003-8045	20030217
JP 2005519079	T2	20050630	JP 2003-570844	20030217
CN 1638766	A	20050713	CN 2003-804767	20030217
US 2003236288	A1	20031225	US 2003-376579	20030228
NO 2004004091	A	20040927	NO 2004-4091	20040927
PRAI DE 2002-10208986	A	20020228		
US 2002-365704P	P	20020319		
WO 2003-EP1560	W	20030217		
OS MARPAT 139:214471				
IT 359848-89-8P				
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)				
(preparation of (alkoxy)(phenyl)oxadiazolones for producing medicaments inhibiting pancreatic lipase)				
RN 359848-89-8 CAPLUS				
CN Urea, N-[4-(5-methoxy-2-oxo-1,3,4-oxadiazol-3(2H)-yl)-2-methylphenyl]-N'-				

[1-(phenylmethyl)-4-piperidinyl] - (9CI) (CA INDEX NAME)



RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2001:676756 CAPLUS

DN 135:242234

TI Preparation of 3-phenyl-5-alkoxy-1,3,4-oxadiazol-2-ones as
hormone-sensitive lipase inhibitors

IN Schoenafinger, Karl; Petry, Stefan; Mueller, Guenter; Baringhaus,
Karl-Heinz

PA Aventis Pharma Deutschland G.m.b.H., Germany

SO PCT Int. Appl., 44 pp.

CODEN: PIXXD2

DT Patent

LA German

FAN.CNT 2

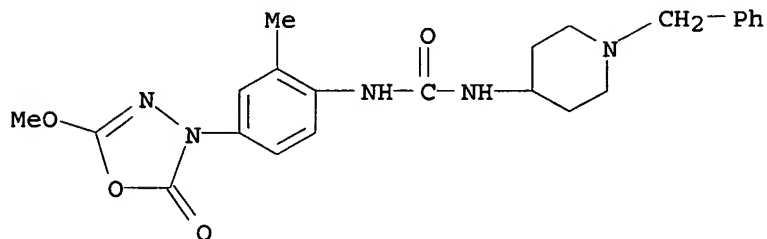
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001066531	A1	20010913	WO 2001-EP1898	20010220
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	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
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	DE 10102265	C1	20020808	DE 2001-10102265	20010118
	CA 2401953	AA	20010913	CA 2001-2401953	20010220
	EP 1263745	A1	20021211	EP 2001-905805	20010220
	EP 1263745	B1	20040519		
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
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	JP 2003525931	T2	20030902	JP 2001-565347	20010220
	EE 200200498	A	20040216	EE 2002-498	20010220
	AT 267184	E	20040615	AT 2001-905805	20010220
	NZ 521207	A	20050429	NZ 2001-521207	20010220
	AU 784827	B2	20060629	AU 2001-33787	20010220
	RU 2281283	C2	20060810	RU 2002-126556	20010220
	NO 2002004201	A	20020903	NO 2002-4201	20020903
PRAI	DE 2000-10010968	A	20000307		
	DE 2001-10102265	A	20010118		
	WO 2001-EP1898	W	20010220		
OS	MARPAT 135:242234				
IT	359848-89-8P				

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use);

BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of 3-phenyl-5-alkoxy-1,3,4-oxadiazol-2-ones as
hormone-sensitive lipase inhibitors)

RN 359848-89-8 CAPLUS

CN Urea, N-[4-(5-methoxy-2-oxo-1,3,4-oxadiazol-3(2H)-yl)-2-methylphenyl]-N'-
[1-(phenylmethyl)-4-piperidinyl]- (9CI) (CA INDEX NAME)



RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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(FILE 'HOME' ENTERED AT 09:35:28 ON 24 AUG 2006)

FILE 'REGISTRY' ENTERED AT 09:35:48 ON 24 AUG 2006

L1 STRUC
L2 50 S L1
L3 18324 S L1 FUL
L4 2850 S L3 AND (PIPERIDIN? OR AZEPIDIN?)
E AZEPIDIN
L5 74 S L3 AND AZEPIN?

FILE 'CAPLUS' ENTERED AT 09:38:59 ON 24 AUG 2006

L6 16 S L5

FILE 'REGISTRY' ENTERED AT 09:40:41 ON 24 AUG 2006

FILE 'CAPLUS' ENTERED AT 09:42:12 ON 24 AUG 2006

FILE 'STNGUIDE' ENTERED AT 09:48:08 ON 24 AUG 2006

FILE 'CAPLUS' ENTERED AT 09:49:53 ON 24 AUG 2006

FILE 'REGISTRY' ENTERED AT 09:50:50 ON 24 AUG 2006

L7 2850 S L4 NOT L5
L8 STRUC
L9 546 SEARCH L8 SSS SUB=L7 FUL
L10 546 SEARCH L8 SSS SUB=L9 FUL

FILE 'CAPLUS' ENTERED AT 09:57:41 ON 24 AUG 2006

L11 2 S L10

FILE 'STNGUIDE' ENTERED AT 09:59:22 ON 24 AUG 2006

FILE 'REGISTRY' ENTERED AT 10:06:36 ON 24 AUG 2006

L12 2304 S L7 NOT L10

FILE 'CAPLUS' ENTERED AT 10:08:33 ON 24 AUG 2006

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44488 PHOSPHOLIPASE

L15 1 L13 AND PHOSPHOLIPASE

=> d bib

L15 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:878375 CAPLUS

DN 141:350047

TI Preparation of phospholipase C inhibitors for use in treating
inflammatory diseases

IN Lagu, Bharat; Rupert, Kenneth; Wachter, Michael

PA Janssen Pharmaceutica N.V., Belg.

SO PCT Int. Appl., 114 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	WO 2004089901	A2	20041021	WO 2004-US9847	20040331
	WO 2004089901	A3	20041209		
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 GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
 LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
 NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
 TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
 RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
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 TD, TG

US 2004242639 A1 20041202 US 2004-814070 20040331
 PRAI US 2003-459078P P 20030331
 OS MARPAT 141:350047

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L15 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:878375 CAPLUS

DN 141:350047

TI Preparation of phospholipase C inhibitors for use in treating
 inflammatory diseases

IN Lagu, Bharat; Rupert, Kenneth; Wachter, Michael

PA Janssen Pharmaceutica N.V., Belg.

SO PCT Int. Appl., 114 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004089901	A2	20041021	WO 2004-US9847	20040331
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US 2004242639 A1 20041202 US 2004-814070 20040331
 PRAI US 2003-459078P P 20030331
 OS MARPAT 141:350047
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* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB This invention is directed to heterocyclyl-substituted anilino
 phospholipase C inhibitor compds. I [X = (un)substituted-amino,
 -heterocyclyl, etc.; R3 = O or S; R4 = cycloalkyl, benzofused dioxolyl,
 benzofused dioxinyl, or aryl; L = a bond or a linking group; R5 =
 (un)substituted-alkyl, -cycloalkyl, or -aryl; Y = (un)substituted-alkyl; n
 = 1-2] useful in treating or ameliorating an inflammatory disorders and/or
 restenosis and enantiomers, diastereomers and pharmaceutically acceptable
 salts thereof. For example, compound II were prepared in a multi-steps
 employing a solid phase synthesis starting from 4-fluoro-3-nitrobenzoic
 acid. The latter inhibits phospholipase C- β 2 with an IC50

= 3.4 μ M.

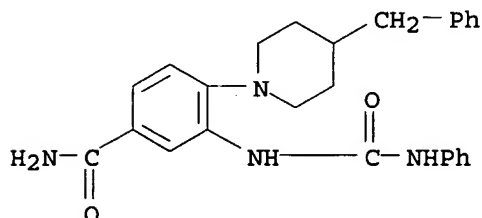
IT 775349-73-0P 775349-74-1P 775349-75-2P
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775349-79-6P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of ureido piperidinyl derivative as phospholipase c inhibitors for treatment of inflammatory disorders)

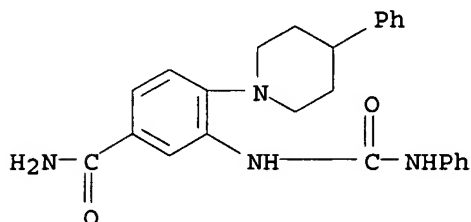
RN 775349-73-0 CAPLUS

CN Benzamide, 3-[[[(phenylamino)carbonyl]amino]-4-[4-(phenylmethyl)-1-piperidinyl]- (9CI) (CA INDEX NAME)



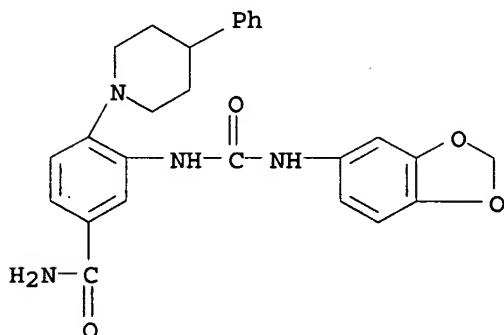
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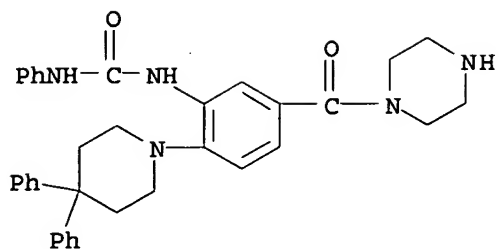
RN 775349-75-2 CAPLUS

CN Benzamide, 3-[[[(1,3-benzodioxol-5-ylamino)carbonyl]amino]-4-(4-phenyl-1-piperidinyl)- (9CI) (CA INDEX NAME)



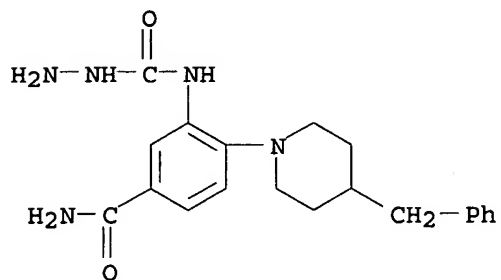
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CN Piperazine, 1-[4-(4,4-diphenyl-1-piperidinyl)-3-[[[(phenylamino)carbonyl]amino]benzoyl]- (9CI) (CA INDEX NAME)



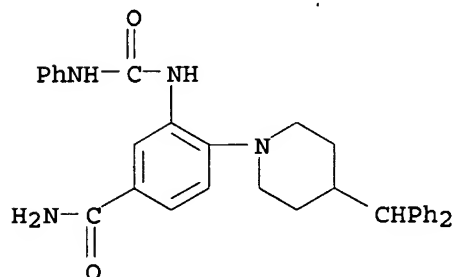
RN 775349-77-4 CAPLUS

CN Hydrazinecarboxamide, N-[5-(aminocarbonyl)-2-[4-(phenylmethyl)-1-piperidinyl]phenyl]- (9CI) (CA INDEX NAME)



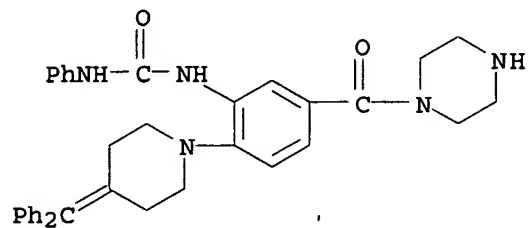
RN 775349-78-5 CAPLUS

CN Benzamide, 4-[4-(diphenylmethyl)-1-piperidinyl]-3-[[(phenylamino)carbonyl]amino]- (9CI) (CA INDEX NAME)

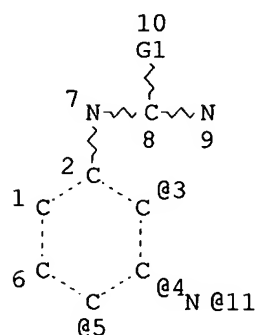


RN 775349-79-6 CAPLUS

CN Piperazine, 1-[4-[4-(diphenylmethylene)-1-piperidinyl]-3-[[(phenylamino)carbonyl]amino]benzoyl]- (9CI) (CA INDEX NAME)



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DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED
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STEREO ATTRIBUTES: NONE
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E AZEPIDIN
L5 74 S L3 AND AZEPIN?
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L6 16 S L5
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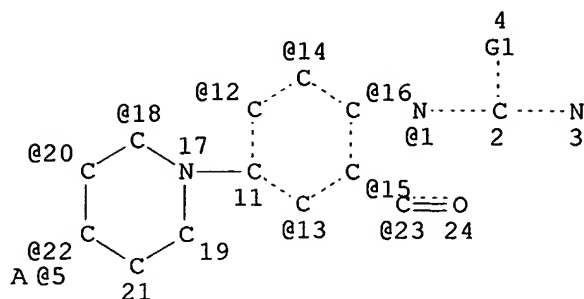
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L8 STRUC
L9 546 SEARCH L8 SSS SUB=L7 FUL
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L8 HAS NO ANSWERS
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L8

STR



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VPA 23-13/15 U

VPA 5-18/20/22 U

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DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RSPEC 22 11

NUMBER OF NODES IS 19

STEREO ATTRIBUTES: NONE

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ENTER SUBSET L# OR (END):19

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100.0% PROCESSED 546 ITERATIONS

546 ANSWERS

SEARCH TIME: 00.00.01

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COST IN U.S. DOLLARS

SINCE FILE TOTAL

ENTRY SESSION

FULL ESTIMATED COST

212.64 467.87

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE TOTAL

ENTRY SESSION

CA SUBSCRIBER PRICE

0.00 -0.75

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FILE COVERS 1907 - 24 Aug 2006 VOL 145 ISS 9
FILE LAST UPDATED: 23 Aug 2006 (20060823/ED)

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<http://www.cas.org/infopolicy.html>

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L11 2 L10

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L11 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:878375 CAPLUS

DN 141:350047

TI Preparation of phospholipase C inhibitors for use in treating inflammatory diseases

IN Lagu, Bharat; Rupert, Kenneth; Wachter, Michael

PA Janssen Pharmaceutica N.V., Belg.

SO PCT Int. Appl., 114 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004089901	A2	20041021	WO 2004-US9847	20040331
	WO 2004089901	A3	20041209		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
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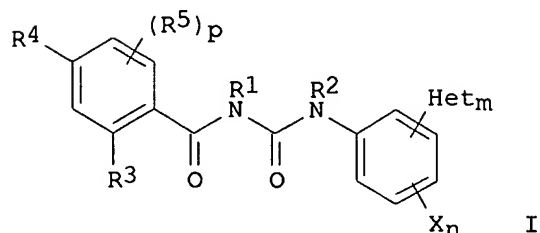
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB This invention is directed to heterocyclyl-substituted anilino phospholipase C inhibitor compds. I [X = (un)substituted-amino, -heterocyclyl, etc.; R3 = O or S; R4 = cycloalkyl, benzofused dioxolyl, benzofused dioxinyl, or aryl; L = a bond or a linking group; R5 = (un)substituted-alkyl, -cycloalkyl, or -aryl; Y = (un)substituted-alkyl; n = 1-2] useful in treating or ameliorating an inflammatory disorders and/or restenosis and enantiomers, diastereomers and pharmaceutically acceptable salts thereof. For example, compound II were prepared in a multi-steps

employing a solid phase synthesis starting from 4-fluoro-3-nitrobenzoic acid. The latter inhibits phospholipase C- β 2 with an IC50 = 3.4 μ M.

L11 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
 AN 2004:60473 CAPLUS
 DN 140:128423
 TI Preparation of heterocyclylbenzoylureas for treating type 2 diabetes
 IN Schoenafinger, Karl; Defossa, Elisabeth; Kadereit, Dieter; Von Roedern, Erich; Klabunde, Thomas; Burger, Hans-Joerg; Herling, Andreas; Wendt, Karl-Ulrich
 PA Aventis Pharma Deutschland GmbH, Germany
 SO PCT Int. Appl., 67 pp.
 CODEN: PIXXD2
 DT Patent
 LA German
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004007455	A1	20040122	WO 2003-EP7078	20030703
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	DE 10320326	A1	20041202	DE 2003-10320326	20030506
	CA 2493374	AA	20040122	CA 2003-2493374	20030703
	AU 2003249937	A1	20040202	AU 2003-249937	20030703
	EP 1523475	A1	20050420	EP 2003-763692	20030703
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	BR 2003012697	A	20050426	BR 2003-12697	20030703
	CN 1668593	A	20050914	CN 2003-816581	20030703
	JP 2006502247	T2	20060119	JP 2005-505058	20030703
	US 2004152743	A1	20040805	US 2003-617498	20030711
	ZA 2004010133	A	20051123	ZA 2004-10133	20041215
	NO 2005000648	A	20050207	NO 2005-648	20050207
PRAI	DE 2002-10231627	A	20020712		
	DE 2003-10306503	A	20030217		
	DE 2003-10320326	A	20030506		
	US 2002-430782P	P	20021204		
	WO 2003-EP7078	W	20030703		
OS	MARPAT 140:128423				
GI					



AB Title compds. [I; R1, R2 = H, (substituted) A, OA, COA, CO2A, AlkCO2H, AlkCO2A; A = alkyl; Alk = alkylene; R3, R4 = F, Cl, Br, OH, NO2, CN, (substituted) A, OA, alkenyloxy, alkynyl; R5 = H, F, Cl, Br, OH, NO2, CN, (substituted) A, OA, COA, AlkCO2H, AlkCO2A, SO2A, alkenyloxy, alkynyl; X = H, F, Cl, Br, OH, NO2, CN, (substituted) A, COA, AlkCO2H, AlkCO2A, SO2A, alkenyl, alkynyl, OA, SO1-2A, NHA, NA2, CO2H, CO2A, CONH2, CONHA, CONA2, SO2NH2, SO2NHA, SO2NA2, NHCOR6; R6 = H, A, cycloalkyl, cycloalkylalkylene, alkenyl, alkynyl, AlkCO2A, AlkCOA, AlkCO2H, AlkCONH2, aryl, Alkaryl, heteroaryl, Alkheteroaryl, heteroarylcarbonyl; het = 4-7 membered (substituted) heterocyclyl, with the exception of pyrrole; m = 1-5; n, p = 0-3], were prepared Thus, 1-(4-amino-3-fluorophenyl)-1H-[1,2,4]triazole (preparation given) and 2-chloro-4,5-difluorobenzoylisocyanate were stirred 30 min in MeCN to give 1-(2-chloro-4,5-difluorobenzoyl)-3-(2-fluor-4-[1,2,4]triazol-1-ylphenyl)urea. The latter at 10 μ M gave 94% inhibition of activated glycogen phosphorylase.

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d hitstr 2

L11 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN

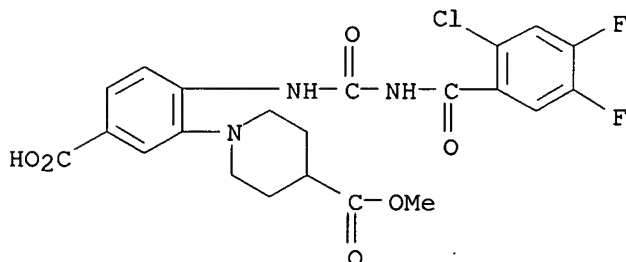
IT 648917-41-3P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

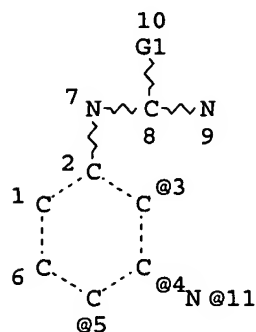
(preparation of heterocyclylbenzoylureas for treating type 2 diabetes)

RN 648917-41-3 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[5-carboxy-2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]phenyl]-, 4-methyl ester (9CI) (CA INDEX NAME)



=> d l1
 L1 HAS NO ANSWERS
 L1 STR



VAR G1=O/S
 VPA 11-3/4/5 U
 NODE ATTRIBUTES:
 NSPEC IS R AT 11
 DEFAULT MLEVEL IS ATOM
 DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
 RSPEC 4
 NUMBER OF NODES IS 11

STEREO ATTRIBUTES: NONE

=> d his l3

(FILE 'REGISTRY' ENTERED AT 09:35:48 ON 24 AUG 2006)
 L3 18324 S L1 FUL

=> d his l5

(FILE 'REGISTRY' ENTERED AT 09:35:48 ON 24 AUG 2006)
 E AZEPIDIN
 L5 74 S L3 AND AZEPIN?

=> s 15

L6 16 L5

=> d bib 1-16

L6 ANSWER 1 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2006:164439 CAPLUS

DN 144:253908

TI Preparation of aryl urea derivatives as CB1 cannabinoid receptor modulators

IN Bloxham, Jason; Fyfe, Matthew Colin Thor; Horswill, James; Jeevaratnam, Revathy Perpetua; Keily, John; Procter, Martin James; Schofield, Karen Lesley; Shaaban, Salam; Swain, Simon Andrew; Wong-Kai-In, Philippe

PA Prosidion Limited, UK

SO PCT Int. Appl., 66 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006018662	A2	20060223	WO 2005-GB50131	20050816
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				

PRAI US 2004-602268P P 20040816

OS MARPAT 144:253908

L6 ANSWER 2 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2005:1242631 CAPLUS

DN 144:6677

TI Preparation of urea containing N-aryl or N-heteroaryl substituted heterocycles as antagonists of P2Y1 receptor for treating thrombotic conditions

IN Tuerdi, Huji; Chao, Hannguang J.; Qiao, Jennifer X.; Wang, Tammy C.; Gungor, Timur

PA USA

SO U.S. Pat. Appl. Publ., 102 pp.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2005261244	A1	20051124	US 2005-126567	20050510
	WO 2005113511	A1	20051201	WO 2005-US16422	20050511
	WO 2005113511	C2	20060202		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK,				

SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU,
 ZA, ZM, ZW
 RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
 AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
 EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT,
 RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,
 MR, NE, SN, TD, TG
 WO 2005113537 A2 20051201 WO 2005-US16525 20050511
 WO 2005113537 A3 20051222
 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
 CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
 GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ,
 LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA,
 NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK,
 SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU,
 ZA, ZM, ZW
 RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
 AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
 EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT,
 RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,
 MR, NE, SN, TD, TG
 PRAI US 2004-570288P P 20040512
 US 2005-665735P P 20050328
 US 2005-665817P P 20050328
 US 2005-126567 A 20050510
 OS MARPAT 144:6677
 L6 ANSWER 3 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
 AN 2003:891946 CAPLUS
 DN 139:381494
 TI Preparation of semicarbazides as inhibitors of blood-coagulation factor Xa
 and VIIa
 IN Mederski, Werner; Tsaklakidis, Christos; Cezanne, Bertram; Dorsch, Dieter;
 Barnes, Christopher; Gleitz, Johannes
 PA Merck Patent Gmbh, Germany
 SO Ger. Offen., 18 pp.
 CODEN: GWXXBX
 DT Patent
 LA German
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 10220048	A1	20031113	DE 2002-10220048	20020504
	CA 2485065	AA	20031113	CA 2003-2485065	20030407
	WO 2003093254	A1	20031113	WO 2003-EP3581	20030407
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2003227569	A1	20031117	AU 2003-227569	20030407
	EP 1501814	A1	20050202	EP 2003-724967	20030407
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	US 2005171102	A1	20050804	US 2003-513451	20030407
	JP 2005530754	T2	20051013	JP 2004-501393	20030407
PRAI	DE 2002-10220048	A	20020504		

WO 2003-EP3581 W 20030407
OS MARPAT 139:381494

L6 ANSWER 4 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2003:154238 CAPLUS
DN 138:204941
TI Preparation of indol-5-ylureas and relate compounds for the treatment of obesity and type II diabetes
IN Schwink, Lothar; Stengelin, Siegfried; Gossel, Matthias
PA Aventis Pharma Deutschland G.m.b.H., Germany
SO PCT Int. Appl., 77 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003015769	A1	20030227	WO 2002-EP8686	20020803
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	DE 10139416	A1	20030306	DE 2001-10139416	20010817
	CA 2457037	AA	20030227	CA 2002-2457037	20020803
	EP 1418906	A1	20040519	EP 2002-774498	20020803
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
	BR 2002011989	A	20040928	BR 2002-11989	20020803
	CN 1555260	A	20041215	CN 2002-818162	20020803
	JP 2005505530	T2	20050224	JP 2003-520728	20020803
	US 2003212070	A1	20031113	US 2002-218034	20020814
	EE 200400055	A	20040415	EE 2004-55	20030803
	NO 2004000678	A	20040513	NO 2004-678	20040216
	ZA 2004001221	A	20041027	ZA 2004-1221	20040216
	US 2004192693	A1	20040930	US 2004-820706	20040409
	US 2004198731	A1	20041007	US 2004-820703	20040409
	US 2004198732	A1	20041007	US 2004-820736	20040409
	US 2004198733	A1	20041007	US 2004-820883	20040409
PRAI	DE 2001-10139416	A	20010817		
	WO 2002-EP8686	W	20020803		
	US 2002-218034	A3	20020814		

OS MARPAT 138:204941

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 5 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676565 CAPLUS
DN 135:247001
TI Oxidation dyeing composition for keratinous fibers and dyeing method using same
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 51 pp.
CODEN: PIXXD2
DT Patent
LA French

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001066072	A1	20010913	WO 2001-FR663	20010306
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	FR 2805738	A1	20010907	FR 2000-2858	20000306
	FR 2805738	B1	20030314		
	CA 2373099	AA	20010913	CA 2001-2373099	20010306
	EP 1181004	A1	20020227	EP 2001-913934	20010306
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	BR 2001005561	A	20020319	BR 2001-5561	20010306
	AU 752948	B2	20021003	AU 2001-39341	20010306
	JP 2003525889	T2	20030902	JP 2001-564725	20010306
	ZA 2001008983	A	20020911	ZA 2001-8983	20011031
	US 2003028977	A1	20030213	US 2002-959702	20020503
PRAI	FR 2000-2858	A	20000306		
	WO 2001-FR663	W	20010306		
OS	MARPAT 135:247001				

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676564 CAPLUS
DN 135:247000
TI Oxidation dyeing composition for keratinous fibers comprising
paraphenylenediamine derivatives and coupling agents
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 56 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001066071	A1	20010913	WO 2001-FR660	20010306
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	FR 2805737	A1	20010907	FR 2000-2857	20000306
	FR 2805737	B1	20030103		
	CA 2373097	AA	20010913	CA 2001-2373097	20010306
	EP 1181005	A1	20020227	EP 2001-915449	20010306
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	BR 2001005562	A	20020319	BR 2001-5562	20010306
	JP 2003525888	T2	20030902	JP 2001-564724	20010306

ZA 2001009069	A	20020613	ZA 2001-9069	20011102
US 2003009835	A1	20030116	US 2002-959704	20020208
US 6890362	B2	20050510		
PRAI FR 2000-2857	A	20000306		
WO 2001-FR660	W	20010306		

OS MARPAT 135:247000

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 7 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676563 CAPLUS
DN 135:246999
TI Oxidation dyeing composition for keratinous fibers containing
paraphenylenediamine derivatives and oxidants
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 44 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001066070	A1	20010913	WO 2001-FR646	20010305
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	FR 2805739	A1	20010907	FR 2000-2860	20000306
	FR 2805739	B1	20030110		
	CA 2400464	AA	20010913	CA 2001-2400464	20010305
	EP 1263399	A1	20021211	EP 2001-911848	20010305
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	BR 2001009175	A	20030422	BR 2001-9175	20010305
	JP 2003525887	T2	20030902	JP 2001-564723	20010305
	US 2003167579	A1	20030911	US 2003-333664	20030410
PRAI	FR 2000-2860	A	20000306		
	WO 2001-FR646	W	20010305		

OS MARPAT 135:246999

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 8 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676562 CAPLUS
DN 135:246998
TI Oxidation dyeing composition for keratinous fibers comprising substituted
paraphenylenediamine derivatives and polymers
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI WO 2001066069 A1 20010913 WO 2001-FR645 20010305
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO,
RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
FR 2805740 A1 20010907 FR 2000-2861 20000306
FR 2805740 B1 20030905
CA 2400459 AA 20010913 CA 2001-2400459 20010305
EP 1263398 A1 20021211 EP 2001-911847 20010305
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
BR 2001009179 A 20030422 BR 2001-9179 20010305
JP 2003528054 T2 20030924 JP 2001-564722 20010305
US 2004088798 A1 20040513 US 2003-363147 20030911
PRAI FR 2000-2861 A 20000306
WO 2001-FR645 W 20010305
OS MARPAT 135:246998
RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676561 CAPLUS
DN 135:246997
TI Oxidation dyeing composition for keratinous fibers with a particular
paraphenylenediamine derivative and a particular direct dyeing agent
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 49 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2001066068	A1	20010913	WO 2001-FR644	20010305
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
FR 2805741	A1	20010907	FR 2000-2862	20000306
FR 2805741	B1	20030620		
CA 2400456	AA	20010913	CA 2001-2400456	20010305
BR 2001009021	A	20021126	BR 2001-9021	20010305
EP 1263397	A1	20021211	EP 2001-911846	20010305
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
JP 2003528053	T2	20030924	JP 2001-564721	20010305
US 2003159221	A1	20030828	US 2003-333663	20030410
PRAI FR 2000-2862	A	20000306		
WO 2001-FR644	W	20010305		
OS MARPAT 135:246997				
RE.CNT 7				

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 10 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2000:646000 CAPLUS

DN 133:222725

TI Preparation of thiazolylureas as antivirals

IN Fischer, Rudiger; Kleymann, Gerald; Baumeister, Judith; Bender, Wolfgang; Betz, Ulrich; Eckenberg, Peter; Handke, Gabriele; Hendrix, Martin; Schneider, Udo; Weber, Olaf; Henninger, Kerstin; Jensen, Axel; Keldenich, Jorg

PA Bayer Aktiengesellschaft, Germany

SO PCT Int. Appl., 133 pp.

CODEN: PIXXD2

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000053591	A1	20000914	WO 2000-EP1498	20000224
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	DE 19959958	A1	20010830	DE 1999-19959958	19991213
	CA 2366607	AA	20000914	CA 2000-2366607	20000224
	EP 1161423	A1	20011212	EP 2000-907614	20000224
	EP 1161423	B1	20041110		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	JP 2002539119	T2	20021119	JP 2000-604030	20000224
	ES 2232427	T3	20050601	ES 2000-907614	20000224
	US 6500817	B1	20021231	US 2001-914554	20010831
PRAI	DE 1999-19910245	A	19990308		
	DE 1999-19959958	A	19991213		
	WO 2000-EP1498	W	20000224		

OS MARPAT 133:222725

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 1998:603272 CAPLUS

DN 129:230732

TI Preparation of N-(2-heterocyclylphenyl)amides as herbicides

IN Andree, Roland; Drewes, Mark Wilhelm; Findeisen, Kurt; Kluth, Joachim; Linker, Karl-Heinz; Mueller, Klaus-Helmut; Schallner, Otto; Dollinger, Markus

PA Bayer A.-G., Germany

SO Ger. Offen., 70 pp.

CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 19708928	A1	19980910	DE 1997-19708928	19970305
	CA 2283298	AA	19980911	CA 1998-2283298	19980220
	WO 9839304	A1	19980911	WO 1998-EP972	19980220

W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG

AU 9866226	A1	19980922	AU 1998-66226	19980220
AU 731129	B2	20010322		
EP 973752	A1	20000126	EP 1998-908103	19980220
EP 973752	B1	20050511		
R: BE, CH, DE, DK, ES, FR, GB, IT, LI, NL				
BR 9808200	A	20000516	BR 1998-8200	19980220
JP 2001513785	T2	20010904	JP 1998-538103	19980220
MX 9908144	A	20000131	MX 1999-8144	19990903
US 6602826	B1	20030805	US 1999-367476	19990920
US 6686318	B1	20040203	US 2003-420203	20030422
PRAI DE 1997-19708928	A	19970305		
WO 1998-EP972	W	19980220		
US 1999-367476	A3	19990920		
OS MARPAT 129:230732				

L6 ANSWER 12 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 1995:758682 CAPLUS

DN 123:169279

TI Preparation of cyclohexanediurea derivatives as ACAT inhibitors

IN Yamada, Toshihiro; Nobuhara, Yoichi; Takagi, Ichinari; Furumoto, Shiho; Kobayashi, Kazuhiro; Ikemoto, Kiyohito

PA Nissin Food Products Co., Ltd., Japan

SO PCT Int. Appl., 146 pp.
CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	WO 9507258	A1	19950316	WO 1994-JP1475	19940907
	W: AU, CA, KR, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	JP 07082232	A2	19950328	JP 1993-226247	19930910
	JP 3286745	B2	20020527		
	CA 2171295	AA	19950316	CA 1994-2171295	19940907
	AU 9476235	A1	19950327	AU 1994-76235	19940907
	AU 680941	B2	19970814		
	EP 718281	A1	19960626	EP 1994-926366	19940907
	EP 718281	B1	20010627		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
	AT 202555	E	20010715	AT 1994-926366	19940907
	US 5733931	A	19980331	US 1996-617828	19960308
PRAI	JP 1993-226247	A	19930910		
	WO 1994-JP1475	W	19940907		
OS	MARPAT 123:169279				

L6 ANSWER 13 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 1995:648096 CAPLUS

DN 123:55933

TI Preparation of 3-phenylureido-1,4-benzodiazepine selective cholecystokinin-B and/or gastrin receptor antagonists

IN Semple, Graeme; Ryder, Hamish; Szelke, Michael; Satoh, Masato; Ohta, Mitsuaki; Miyata, Keiji; Nishida, Akito

PA Yamanouchi Pharmaceutical Co., Ltd., Japan; Ferring Research Ltd.

SO PCT Int. Appl., 20 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9506041	A1	19950302	WO 1994-GB1858	19940825
	W: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, JP, KE, KG, KP, KR, KZ, LK, LT, LU, LV, MD, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TJ, TT, UA, US, UZ, VN				
	RW: KE, MW, SD, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	GB 2282594	A1	19950412	GB 1993-17692	19930825
	AU 9474660	A1	19950321	AU 1994-74660	19940825
PRAI	GB 1993-17692	A	19930825		
	WO 1994-GB1858	W	19940825		
OS	MARPAT 123:55933				

L6 ANSWER 14 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 1992:531008 CAPLUS

DN 117:131008

TI Amidoxime functions as neighboring groups in cyclodehydrogenation reactions

AU Moehrle, Hans; Lessel, Juergen

CS Inst. Pharm. Chem., Univ. Duesseldorf, Duesseldorf, W-4000/1, Germany

SO Chemische Berichte (1992), 125(8), 1843-9

CODEN: CHBEAM; ISSN: 0009-2940

DT Journal

LA German

OS CASREACT 117:131008

L6 ANSWER 15 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 1978:563531 CAPLUS

DN 89:163531

TI Cyclodehydrogenation of aniline derivatives with aromatic carboxamide or urea functions

AU Moehrle, Hans; Hemmerling, Hans Joerg

CS Inst. Pharm. Chem., Univ. Duesseldorf, Duesseldorf, Fed. Rep. Ger.

SO Archiv der Pharmazie (Weinheim, Germany) (1978), 311(7), 586-94

CODEN: ARPMAS; ISSN: 0365-6233

DT Journal

LA German

OS CASREACT 89:163531

L6 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 1975:156229 CAPLUS

DN 82:156229

TI Heterocyclic syntheses. XXVIII. ortho-(Dialkylamino)benzenesulfonyl azides, -benzoyl azides, and -benzoyldiazomethanes

AU Martin, John; Meth-Cohn, Otto; Suschitzky, Hans

CS Ramage Lab., Univ. Salford, Salford, UK

SO Journal of the Chemical Society, Perkin Transactions 1: Organic and Bio-Organic Chemistry (1972-1999) (1974), (21), 2451-8

CODEN: JCPRB4; ISSN: 0300-922X

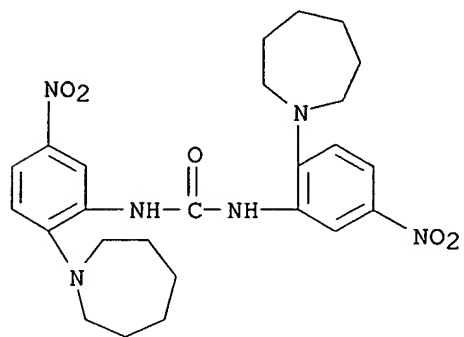
DT Journal

LA English

OS CASREACT 82:156229

=> d hitstr 16

L6 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 55259-36-4P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 55259-36-4 CAPLUS
CN Urea, N,N'-bis[2-(hexahydro-1H-azepin-1-yl)-5-nitrophenyl]- (9CI) (CA
INDEX NAME)



(FILE 'CAPLUS' ENTERED AT 09:38:59 ON 24 AUG 2006)
L6 16 S L5

FILE 'REGISTRY' ENTERED AT 09:40:41 ON 24 AUG 2006

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YOU HAVE REQUESTED DATA FROM FILE 'CAPLUS' - CONTINUE? (Y)/N:end

=> fil caplus

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
1.32	206.50

FULL ESTIMATED COST

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FILE LAST UPDATED: 23 Aug 2006 (20060823/ED)

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<http://www.cas.org/infopolicy.html>

=> d hitstr 15

L6 ANSWER 15 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

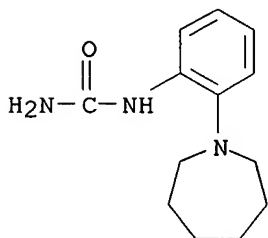
IT 67829-57-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and cyclodehydrogenation of)

RN 67829-57-6 CAPLUS

CN Urea, [2-(hexahydro-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



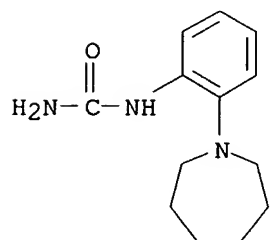
IT 67829-58-7P

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

RN 67829-58-7 CAPLUS
CN Urea, [2-(hexahydro-1H-azepin-1-yl)phenyl]-, monoperchlorate (9CI) (CA INDEX NAME)

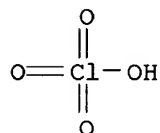
CM 1

CRN 67829-57-6
CMF C13 H19 N3 O



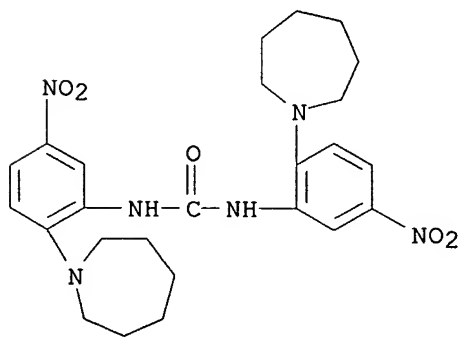
CM 2

CRN 7601-90-3
CMF Cl H O4



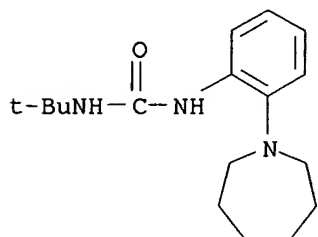
=> d hitstr 16

L6 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 55259-36-4P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 55259-36-4 CAPLUS
CN Urea, N,N'-bis[2-(hexahydro-1H-azepin-1-yl)-5-nitrophenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 14

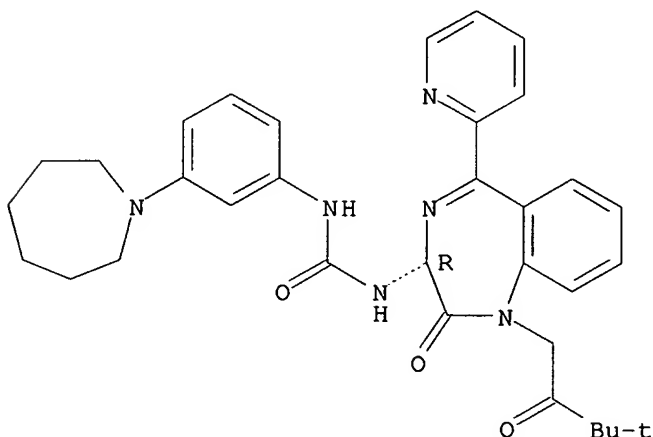
L6 ANSWER 14 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 140410-89-5P
RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
(preparation and spectra of)
RN 140410-89-5 CAPLUS
CN Urea, N-(1,1-dimethylethyl)-N'-[2-(hexahydro-1H-azepin-1-yl)phenyl]- (9CI)
(CA INDEX NAME)



=> d hitstr 13

L6 ANSWER 13 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 164343-30-0P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(claimed compound; preparation of 3-phenylureido-1,4-benzodiazepine selective cholecystinin-B and/or gastrin receptor antagonists)
RN 164343-30-0 CAPLUS
CN Urea, N-[1-(3,3-dimethyl-2-oxobutyl)-2,3-dihydro-2-oxo-5-(2-pyridinyl)-1H-1,4-benzodiazepin-3-yl]-N'-[3-(hexahydro-1H-azepin-1-yl)phenyl]-, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



=> d hitstr 12

L6 ANSWER 12 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

IT 166967-92-6P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses).

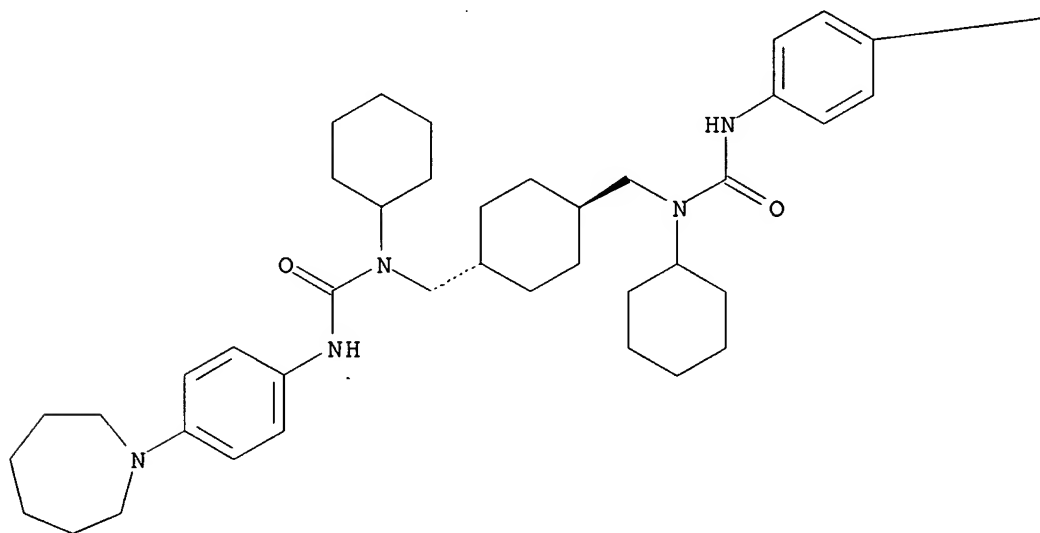
(preparation of cyclohexanediurea derivs. as ACAT inhibitors)

RN 166967-92-6 CAPLUS

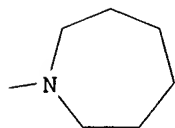
CN Urea, N,N''-[1,4-cyclohexanediylbis(methylene)]bis[N-cyclohexyl-N'-[4-(hexahydro-1H-azepin-1-yl)phenyl]-, dihydrochloride, trans- (9CI) (CA INDEX NAME)

Relative stereochemistry.

PAGE 1-A



PAGE 1-B



●2 HCl

=> d hitstr 11

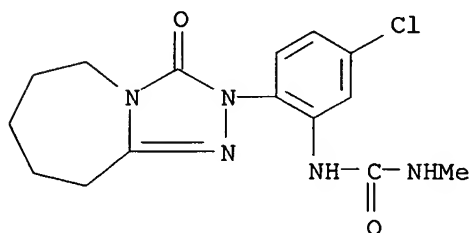
L6 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

IT 212903-67-8P 212903-72-5P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of N-(2-heterocyclylphenyl)amides as herbicides)

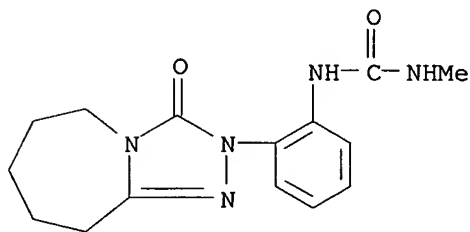
RN 212903-67-8 CAPLUS

CN Urea, N-[5-chloro-2-(6,7,8,9-tetrahydro-3-oxo-3H-1,2,4-triazolo[4,3-a]azepin-2(5H)-yl)phenyl]-N'-methyl- (9CI) (CA INDEX NAME)



RN 212903-72-5 CAPLUS

CN Urea, N-methyl-N'-[2-(6,7,8,9-tetrahydro-3-oxo-3H-1,2,4-triazolo[4,3-a]azepin-2(5H)-yl)phenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 10

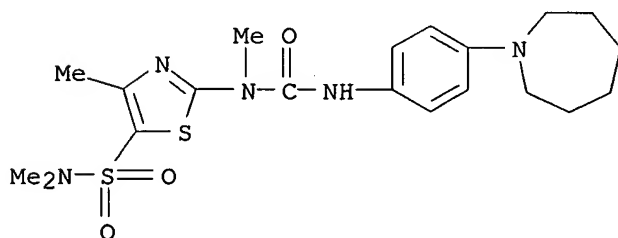
L6 ANSWER 10 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

IT 292137-30-5P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of thiazolylureas as antivirals)

RN 292137-30-5 CAPLUS

CN 5-Thiazolesulfonamide, 2-[[[4-(hexahydro-1H-azepin-1-yl)phenyl]amino]carbonyl]methylamino]-N,N,4-trimethyl- (9CI) (CA INDEX NAME)



=> d hitstr 9

L6 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

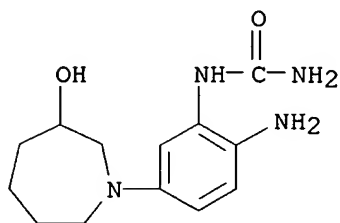
IT 359841-52-4

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
(Uses)

(oxidative hair dyes containing paraphenylenediamine derivs. direct dyes)

RN 359841-52-4 CAPLUS

CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA
INDEX NAME)



=> d hitstr 8

L6 ANSWER 8 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

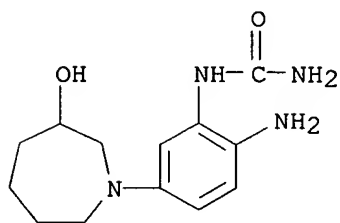
IT 359841-52-4

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
(Uses)

(oxidative hair dyes comprising substituted paraphenylenediamine
derivs. and polymers)

RN 359841-52-4 CAPLUS

CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA
INDEX NAME)



=> d hitstr 7

L6 ANSWER 7 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

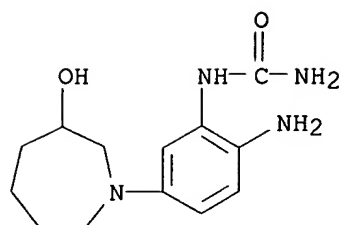
IT 359841-52-4

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
(Uses)

(oxidation dyeing composition for keratinous fibers containing
paraphenylenediamine
derivs. and oxidants)

RN 359841-52-4 CAPLUS

CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA
INDEX NAME)



=> d hitstr 6

L6 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

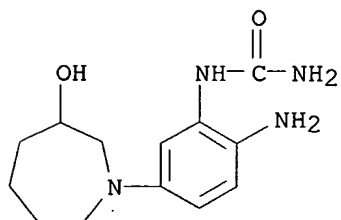
IT 359841-52-4

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

(oxidation dyeing composition for keratinous fibers comprising paraphenylenediamine derivs. and coupling agents)

RN 359841-52-4 CAPLUS

CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 5

L6 ANSWER 5 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

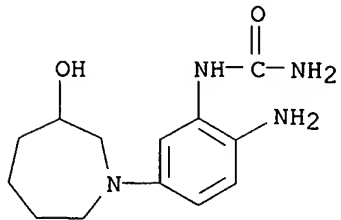
IT 359841-52-4

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

(oxidative hair dye preparation containing paraphenylenediamine derivs.)

RN 359841-52-4 CAPLUS

CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 4

L6 ANSWER 4 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

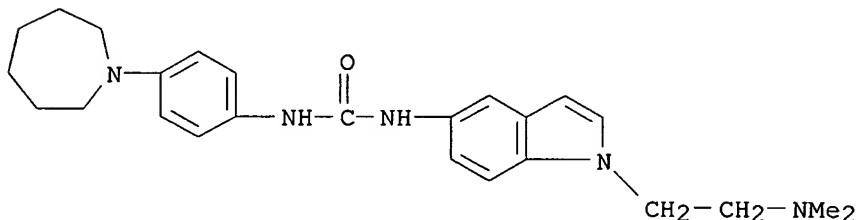
IT 500013-85-4P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of indolylureas and relate compds. for the treatment of obesity and type II diabetes)

RN 500013-85-4 CAPLUS

CN Urea, N-[1-[2-(dimethylamino)ethyl]-1H-indol-5-yl]-N'-[4-(hexahydro-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 3

L6 ANSWER 3 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

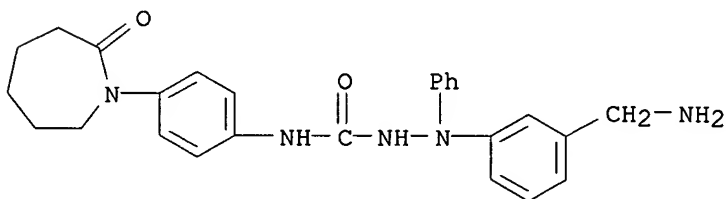
IT 623934-46-3P 623934-47-4P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of semicarbazides as inhibitors of blood-coagulation factor Xa and VIIa)

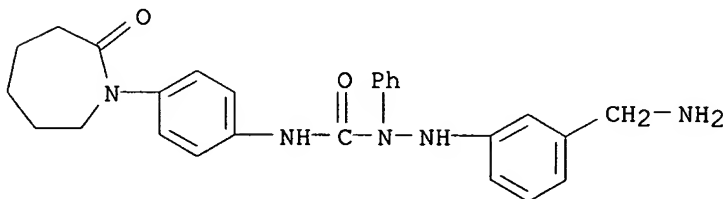
RN 623934-46-3 CAPLUS

CN Hydrazinecarboxamide, 2-[3-(aminomethyl)phenyl]-N-[4-(hexahydro-2-oxo-1H-azepin-1-yl)phenyl]-2-phenyl- (9CI) (CA INDEX NAME)



RN 623934-47-4 CAPLUS

CN Hydrazinecarboxamide, 2-[3-(aminomethyl)phenyl]-N-[4-(hexahydro-2-oxo-1H-azepin-1-yl)phenyl]-1-phenyl- (9CI) (CA INDEX NAME)



=> d hitstr 2

L6 ANSWER 2 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

IT 870072-15-4P

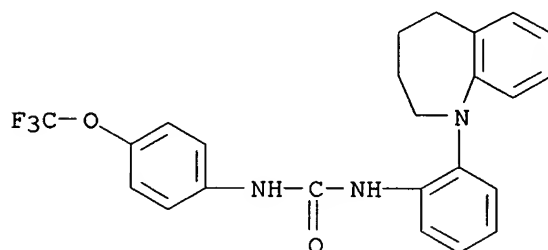
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of urea containing N-aryl or N-heteroaryl substituted heterocycles

as antagonists of P2Y1 receptor for treating thrombotic conditions)

RN 870072-15-4 CAPLUS

CN Urea, N-[2-(2,3,4,5-tetrahydro-1H-1-benzazepin-1-yl)phenyl]-N'-[4-(trifluoromethoxy)phenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 1

'HITSTR' IS NOT A VALID FORMAT FOR FILE 'CAPLUS'

The following are valid formats:

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CBIB ----- AN, plus Compressed Bibliographic Data
CLASS ----- IPC, NCL, ECLA, FTERM
DALL ----- ALL, delimited (end of each field identified)
DMAX ----- MAX, delimited for post-processing
FAM ----- AN, PI and PRAI in table, plus Patent Family data
FBIB ----- AN, BIB, plus Patent FAM
IND ----- Indexing data
IPC ----- International Patent Classifications
MAX ----- ALL, plus Patent FAM, RE
PATS ----- PI, SO
SAM ----- CC, SX, TI, ST, IT
SCAN ----- CC, SX, TI, ST, IT (random display, no answer numbers;
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IABS ----- ABS, indented with text labels
IALL ----- ALL, indented with text labels
IBIB ----- BIB, indented with text labels
IMAX ----- MAX, indented with text labels
ISTD ----- STD, indented with text labels

OBIB ----- AN, plus Bibliographic Data (original)
 OIBIB ----- OBIB, indented with text labels

SBIB ----- BIB, no citations
 SIBIB ----- IBIB, no citations

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 HITSTR ----- HIT RN, its text modification, its CA index name, and
 its structure diagram
 HITSEQ ----- HIT RN, its text modification, its CA index name, its
 structure diagram, plus NTE and SEQ fields
 FHITSTR ----- First HIT RN, its text modification, its CA index name, and
 its structure diagram
 FHITSEQ ----- First HIT RN, its text modification, its CA index name, its
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L6 ANSWER 1 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
 IT 877202-75-0P, 1-[4-(Azepan-1-yl)-3-fluorophenyl]-3-(4-chlorophenyl)urea
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (drug candidate; preparation of aryl urea derivs. as CB1 receptor modulators)
 RN 877202-75-0 CAPLUS
 CN Urea, N-(4-chlorophenyl)-N'-[3-fluoro-4-(hexahydro-1H-azepin-1-yl)phenyl]-
 (9CI) (CA INDEX NAME)

